



Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

JAN 29 2001

UNIX Electronic Co., LTD
c/o Mr. Guenter Ginsberg
Media Trade Corporation
11820 Red Hibiscus Drive
Bonita Springs, FL 34135

Re: K002791

Trade Name: Unix Wrist Digital Blood Pressure Monitor, Model UBP-101S/ UBP-102S
Regulatory Class: II (two)
Product Code: DXN
Dated: December 15, 2000
Received: December 18, 2000

Dear Mr. Ginsberg:

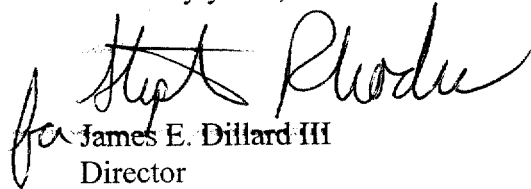
We have reviewed your Section 510(k) notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. A substantially equivalent determination assumes compliance with the Current Good Manufacturing Practice requirements, as set forth in the Quality System Regulation (QS) for Medical Devices: General regulation (21 CFR Part 820) and that, through periodic QS inspections, the Food and Drug Administration (FDA) will verify such assumptions. Failure to comply with the GMP regulation may result in regulatory action. In addition, FDA may publish further announcements concerning your device in the Federal Register. Please note: this response to your premarket notification submission does not affect any obligation you might have under sections 531 through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal laws or regulations.

This letter will allow you to begin marketing your device as described in your 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4648. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its internet address "<http://www.fda.gov/cdrh/dsma/dsmamain.html>".

Sincerely yours,

A handwritten signature in black ink, appearing to read "James E. Dillard III", is written over a pre-printed name. To the left of the signature is a small, stylized handwritten mark that looks like "fa".

James E. Dillard III

Director

Division of Cardiovascular and
Respiratory Devices

Office of Device Evaluation

Center for Devices and

Radiological Health

510(k) NUMBER (IF KNOWN): K002791

DEVICE NAME: UNIX WRIST DIGITAL BLOOD PRESSURE MONITOR

INDICATIONS FOR USE:

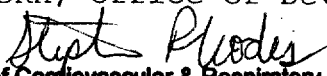
Indication for Use Statement

Device Name: Unix Wrist Digital Blood Pressure Monitor
Model UBP-101S/UBP-102S

The UNIX Blood Pressure Monitor is to be used by adults as a Home Health Care device to monitor Blood Pressure (systolic and diastolic) and the Pulse rate at home. The Blood Pressure Measurement is obtained by the use of the *oscillometric method*, meaning that the monitor detects the blood's movement through the artery in the wrist and converts the movements into digital readings. The Device is not intended to be used by people who have circulatory problems, such as arrhythmia, or if pregnant.

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED.)

Concurrence of CDRH, Office of Device Evaluation (ODE)


Division of Cardiovascular & Respiratory Devices
510(k) Number K002791

Prescription Use _____
(Per 21 CFR 801.109)

OR

Over-The-Counter-Use X
(Optional Format 1-2-96)